

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : A61K 7/00, 7/48, 7/15		A1	(11) International Publication Number: WO 93/25177 (43) International Publication Date: 23 December 1993 (23.12.93)
(21) International Application Number: PCT/US93/05455 (22) International Filing Date: 9 June 1993 (09.06.93)		(81) Designated States: AU, BR, CA, FI, NO, NZ, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(30) Priority data: 07/900,219 17 June 1992 (17.06.92) US		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(71) Applicant: THE PROCTER & GAMBLE COMPANY [US/US]; One Procter & Gamble Plaza, Cincinnati, OH 45202 (US).			
(72) Inventor: LUKE, William, Michael ; 9220 Deer Cross Parkway, 2A, Blue Ash, OH 45236 (US).			
(74) Agents: REED, T., David et al.; The Procter & Gamble Company, 5299 Spring Grove Avenue, Cincinnati, OH 45217 (US).			
(54) Title: COOLANT COMPOSITIONS WITH REDUCED STINGING			
(57) Abstract			
<p>The present invention provides coolant compositions for topical application to the skin which can provide long-lasting cooling with low or reduced skin sting. The compositions comprise a mixture of specific type of acyclic carboxamide coolant with a specific type of 3-substituted-p-menthane coolant, an aqueous vehicle, and no more than about 30 % C₁-C₆ monohydric alcohol.</p>			

CA 002446

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	CA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brasil	IT	Italy	RO	Rumania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Mosocco	TC	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam

CA 002447

- 1 -

COOLANT COMPOSITIONS WITH REDUCED STINGING

TECHNICAL FIELD

This invention relates to low-stinging coolant compositions
5 for application to the skin characterized by containing a multi-
plicity of specific coolant compounds and low or zero levels of
alcohol solvents.

BACKGROUND

Compositions of various types have incorporated within them
10 components which provide cooling sensation to skin. Such compo-
sitions include perfumes, lotions, shaving cream and gels, post-
shaving preparations, shampoos, antiperspirants, deodorants,
anti-acne medicines, first aid ointments, and a variety of other
15 skin care and pharmaceutical products that are applied to the
skin.

It is well established that the "cooling" effect of menthol
is a physiological effect due to the direct action of menthol on
the nerve endings of the human body responsive for the detection
of hot or cold and is not due to latent heat of evaporation. It
20 is believed that the menthol acts as a direct stimulus on the cold
receptors at the nerve endings which in turn stimulate the central
nervous system.

Although menthol is well established as a physiological
coolant, its use and effectiveness is limited in some types of
25 compositions by its strong minty odor and its relative volatility,
particularly for perfumes, colognes, pre- or post-shave lotions,
etc., where aroma is an important aesthetic attribute. The high
volatility of menthol limits the period of time which it can
provide cooling sensation. The high volatility of menthol can
30 also result in eye sting for compositions applied to the face or
in the vicinity of the eyes.

The cooling effect of menthol and other related terpene
alcohols and their derivatives has also been studied and reported
in Koryo, 95, (1970), pp. 39-43. 2,3-p-menthane diol has also been

- 2 -

reported as having a sharp cooling taste (Beilstein, Handbuch der Organischen Chemie, 4th Ed. (1923) Vol. 6, p. 744).

Carboxamides have also been disclosed for use as coolants in a variety of compositions. Two patents describing such materials and compositions are U.S. 4,136,163, January 23, 1979 to Watson et al. and U.S. 4,230,688, October 28, 1980 to Rowsell et al. These patents as well as those set forth above are incorporated herein in their entirety by reference.

Ethanol and other volatile alcohols are commonly used in topical skin compositions to provide a coolant effect. The cooling sensation from volatile alcohols is due primarily to the latent heat of evaporation from the skin. After shave and pre-shave lotions typically contain high levels of ethanol, since it provides a cooling effect and does not interfere with the pleasing aroma typically associated with these products. Ethanol provides a strong initial cooling sensation, however cooling dissipates relatively rapidly as the ethanol evaporates. Unfortunately, the volatile alcohols also tend to impart a stinging sensation in addition to cooling, particularly with respect to skin that has been cut or recently shaved.

It is an object of this invention to provide improved coolant compositions for topical application to the skin.

In particular, it is an object of this invention to provide coolant compositions for application to the skin which can provide an improved cooling in combination with low sting to the skin and eyes.

It is a further object of this invention to provide coolant compositions, as described above, which can provide improved cooling over an extended period of time.

It is yet another object of this invention to provide such improved coolant compositions which need not contain aromatic ingredients, such as menthol, which can cause eye sting or adversely interfere with other fragrances incorporated into the composition.

- 3 -

The invention hereof, including its essential elements as well as a variety of additional ingredients, is described below. The compositions and method hereof can comprise, consist of, or consist essentially of the essential elements as well as any of 5 the additional ingredients or limitations discussed herein.

All percentages reported herein are by weight of the total composition and all ratios are by weight unless otherwise specifically indicated.

SUMMARY OF THE INVENTION

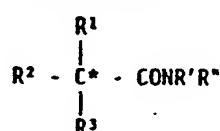
10 The present invention provides aqueous coolant compositions useful for topical application to the skin which impart minimal or no sting and are characterized by both good initial cooling and long lasting cooling sensation upon application to the skin, without requiring the use of odoriferous coolants. The compositions hereof are particularly suitable for pre- and post-shave 15 lotions, perfumes, colognes, etc.

The compositions hereof comprise mixture of particular coolants in an aqueous carrier which contains no more than about 30%, by weight, C₁-C₆ alcohol.

20 The present invention also provides a method for providing a cooling sensation to the skin comprising applying an effective amount of a composition hereof to the skin.

More specifically, the compositions of the present invention comprise:

25 (I) a first coolant component which is an acyclic carboxamide coolant component of the formula:



30

where:

(i) R' and R'' independently are hydrogen, C₁-C₃ alkyl, or C₁-C₃ hydroxyalkyl, R' and R'' provide a total of no more than 8 carbon atoms, and when R' is hydrogen, R'' may also be alkyl-carboxyalkyl of up to 6 carbon atoms

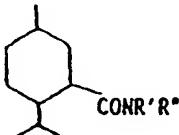
- 4 -

(ii) R' and R'', when taken together, represent an alkylene group of up to 6 carbon atoms, the opposite ends of which are attached to the amide nitrogen atom to form a nitrogen heterocycle, the carbon chain of which optionally being interrupted by oxygen;

5 (iii) R' is hydrogen or C₁-C₅ alkyl; R² and R³ independently are C₁-C₅ alkyl; with the proviso that: (a) R¹, R², and R³ together provide a total of at least 5 carbon atoms; and (b) when R¹ is hydrogen, R² is C₂-C₅ alkyl and R³ is C₃-C₅ alkyl and at least one of R² and R³ is branched;

10 (II) a second coolant component which is a 3-substituted-p-menthane of the formula:

15



where

R' is hydrogen or an aliphatic radical containing up to 25 carbon atoms;

20 R'' is hydroxy or an aliphatic radical containing up to 25 carbon atoms, with the proviso that when R' is hydrogen R'' may also be an aryl radical of up to 10 carbon atoms selected from the group consisting of substituted phenyl, phenalkyl and substituted naphthyl, and pyridyl; and

25 R' and R'', when taken together with the nitrogen atom to which they are attached, represent a cyclic or heterocyclic group of up to 25 carbon atoms;

(III) an aqueous vehicle suitable for topical application to the skin; and

30 (IV) a cosmetically active ingredient or medicament;

wherein said composition contains no more than about 30% by weight of C₁-C₆ alcohol.

35

- 5 -

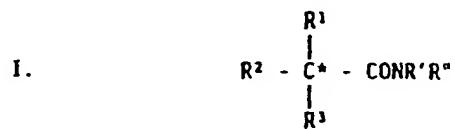
In other embodiments hereof, the compositions contain other cosmetic ingredients or medicaments suitable for application to the skin in place of or in addition to fragrance.

DETAILED DESCRIPTION OF THE INVENTION

5 The essential as well as various optional elements of the compositions of the invention are described below. In general, the compositions contain a mixture of particular carboxamide coolants, water, and no more than about 30%, by weight, of C₁-C₆ monohydric alcohols. The compositions hereof also contain a
10 perfume or other cosmetic ingredient or medicament suitable for application to the skin, or a mixture thereof.

First Coolant Component

The compositions hereof contain as an essential component certain acyclic tertiary and/or secondary carboxamides of the
15 following formula:



20 where R' and R'', when taken separately, are each hydrogen, C₁-C₅ alkyl or C₁-C₆ hydroxyalkyl and provide a total of no more than 8 carbon atoms, with the proviso that when R' is hydrogen R'' may also be alkylcarboxyalkyl of up to 6 carbon atoms;

25 R' and R'', when taken together, represent an alkylene group of up to 6 carbon atoms, the opposite ends of which group are attached to the amide nitrogen atom thereby to form a nitrogen heterocycle, the carbon chain of which may optionally be interrupted by oxygen;

30 R¹ is hydrogen or C₁-C₅ alkyl; and R² and R³ are each C₁-C₅ alkyl; with the provisos that (i) R¹, R² and R³ together provide a total of at least 5 carbon atoms, preferably from 5-10 carbon atoms; and (ii) when R¹ is hydrogen, R² is C₁-C₅ alkyl and R³ is C₃-C₅ alkyl and at least one of R² and R³ is branched, preferably in an alpha or beta position relative to the carbon atom marked
35 (*) in the formula.

- 6 -

The preferred amides used in this invention are the tertiary compounds, i.e., those where each of R¹, R² and R³ is C₁-C₅ alkyl, especially those where R¹ is methyl, ethyl or n-propyl and at least one of R² and R³ is a branched chain group having branching in an alpha or beta position relative to the C atom marked (*) in the formula. Also preferred are non-substituted amides, i.e., where R' is H, and disubstituted amides where R' and R'' are methyl or ethyl. A further preferred group consist of amides of the formula given where R¹ is hydrogen and at least one of R² and R³ is branched in an alpha position relative to the carbon atom marked * in the formula.

Other preferred coolants within the scope of the above formula are monosubstituted tertiary amides of the formula:



where R¹, R² and R³ are each C₁-C₅ alkyl and together provide a total of at least 5, preferably 5-10 carbon atoms; and R' is C₁-C₅ alkyl, C₁-C₈ hydroxyalkyl or alkylcarboxyalkyl of up to 8 carbon atoms. In this group R¹ is preferably methyl, ethyl or n-propyl and one or both of R² and R³ is branched in an alpha or beta position relative to the carbon atom marked (*).

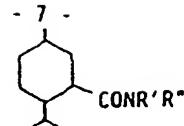
Coolants of the above description are also described in U.S. Patent 4,230,688, Rowsell et al., issued October 28, 1990 (Wilkinson Sword Limited, England), which is incorporated herein by reference.

Second Coolant Component

The compositions hereof contain as a second essential coolant component an N-substituted-p-menthane-3-carboxamide, or a particular type of ketal as will be described below, or a combination thereof.

The N-substituted-p-menthane-3-carboxamides are 3-substituted-p-menthanes of the formula:

II



where R', when taken separately, is hydrogen or an aliphatic radical containing up to 25 carbon atoms; R" when taken separately is hydroxy, or an aliphatic radical containing up to 25 carbon atoms, with the proviso that when R' is hydrogen R" may also be an aryl radical of up to 10 carbon atoms and selected from the group consisting of substituted phenyl, phenalkyl or substituted phenalkyl, naphthyl and substituted naphthyl, pyridyl; and
R' and R", when taken together with the nitrogen atom to which they are attached, represent a cyclic or heterocyclic group of up to 25 carbon atoms, e.g. piperidino, morpholino etc.

In the above definitions "aliphatic" is intended to include any straight-chained, branched-chained or cyclic radical free or aromatic unsaturation, and thus embraces alkyl, cycloalkyl, alkenyl, cycloalkenyl, alkynyl, hydroxylalkyl, acyloxylalkyl, alkoxy, alkoxyalkyl, aminoalkyl, acylaminoalkyl, carboxylalkyl and similar combinations.

Typical values for R' and R" when aliphatic are methyl, ethyl, propyl, butyl, isobutyl, n-decyl, cyclopropyl, cyclohexyl, cyclopentyl, cycloheptylmethyl, 2-hydroxyethyl, 3-hydroxy-n-propyl, 6-hydroxy-n-hexyl, 2-aminoethyl, 2-acetoxyethyl, 2-ethylcarboxyethyl, 4-hydroxybut-2-ynyl, carboxymethyl etc.

When R" is aryl typical values are benzyl, naphthyl, 4-methoxyphenyl, 4-hydroxyphenyl, 4-methylphenyl, 3-hydroxy-4-methylphenyl, 4-fluorophenyl, 4-nitrophenyl, 2-hydroxynaphthyl, pyridyl, etc.

Substitution of the amide group in the 3-position of the p-menthane structure also gives rise to optical isomerism, each of the above-mentioned four geometric isomers, existing in d, l and dl forms. The physiological cooling effect is believed to generally be greater in the l-form than in d-form.

When either R' and R" is aliphatic the preferred values are C₁-C₆, straight or branched chain alkyl, C₁-C₆, straight or branched

- 8 -

chain hydroxyalkyl or aminoalkyl and C₁-C₄ acylated derivatives thereof, and -C_nH_{2n}COR''' or -C_nH_{2n}COOR''', where -C_nH_{2n} is a straight or branched chain alkylene radical in which n is an integer of from 1-6 and R''' is hydrogen or a C₁-C₄ alkyl or hydroxyalkyl group, preferably a C₁-C₄ straight chain alkyl group.

More preferred are monosubstituted compounds, i.e., where R' is H. Also preferred are di-substituted compounds where R' and R" are both C₁-C₄ alkyl also show a very pronounced cooling effect. Most preferred of all are compounds where R' is H and R" is C₁-C₄ alkyl, C₁-C₄ hydroxyalkyl, or -CH₂COOR''', where R''' is C₁-C₄ alkyl.

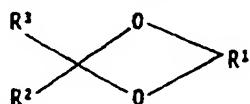
Also included within the scope of this invention are compounds where R' is H and R" is hydroxy or substituted phenyl, e.g., alkylphenyl, hydroxyphenyl, alkoxyphenyl, halophenyl of up to 10 carbon atoms, phenalkyl or substituted phenalkyl e.g., benzyl, naphthyl or substituted naphthyl, and compounds where R' and R" are joined to form a cyclic group. When so joined R' and R" preferably represent an alkylene chain, optionally interrupted by oxygen, which together with the nitrogen atom to which R' and R" are attached forms a 5- or 6-membered heterocyclic ring.

Compounds of the type described above are described in U.S. Patent 4,136,163, Watson et al., issued January 23, 1979 (Wilkinson Sword Limited, England), which is incorporated herein by reference.

Optionally, other coolant ingredients can be included in the compositions hereof to benefit from the particular characteristics of such coolants. Any coolant ingredient suitable for topical application to skin can be used. One type which is particularly preferred includes ketal coolant compounds the formula:

30

III



in which R¹ represents a C₂-C₆-alkylene radical having at least 1,

35

- 9 -

but not more than 3, hydroxyl group(s), preferably 1 hydroxyl group, and either R² and R³ independently of one another represent C₁-C₁₀-alkyl which is optionally substituted by 1 to 3 radicals selected from the group comprising hydroxyl, amino and halogen (such as fluorine, chlorine, bromine or iodine), C₃-C₇-cycloalkyl, preferably cyclohexyl, C₆-C₁₂-aryl, preferably phenyl, with the proviso that the total of the carbon atoms of R² and R³ is not less than 3, or R² and R³ together represent an alkylene radical which, together with the carbon atom which carries the radicals R² and R³, forms a 5-7-membered ring, it being possible for this alkylene radical, in turn, to be substituted by C₁-C₆-alkyl groups.

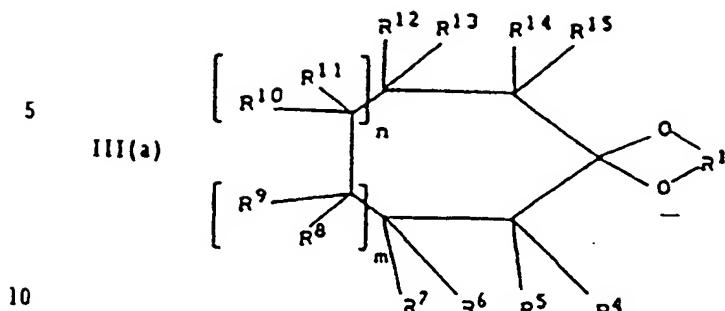
Preferred radicals R² and R³ comprise methyl, isopropyl and tert.-butyl.

The length of the radicals R² and R³ influences the effect of the compounds: shorter radicals lead to an immediate, short effect; longer radicals lead to a delayed, but prolonged effect. An important aspect for the cosmetics industry is the solubility of the compounds in water; this is the case, in particular, with short radicals R² and R³. However if prolonged cooling effect is desired, the water insoluble ketal coolants can be emulsified (including microemulsions) or otherwise dispersed in the compositions hereof through conventional techniques, such as with emulsion-forming surfactants, suspending agents, or solubilization in solvent which is then dispersed, emulsified, or suspended in the composition.

Preferred radicals R¹ embrace 1,2- and 1,3-alkylene radicals which, together with the two oxygen atoms and with the carbon atom to which the two oxygen atoms are attached, form a dioxolane or dioxane ring.

Preferred compounds III in which R² and R³ together represent an alkylene radical are those of the formula

- 10 -



in which R⁴ to R¹⁵ independently of one another denote hydrogen or C₁-C₆-alkyl, preferably hydrogen or C₁-C₄-alkyl, and m and n independently of one another denote zero or 1.

15 Preferred compounds of the formula III(a), are those in which the total of $m + n$ is 1, i.e. ketals of an optionally substituted cyclohexanone.

Preferred substituents, of which there may be present, in particular, 1 to 3, are methyl, isopropyl and tert.-butyl.

20 The ketals III can be prepared by known processes. For example, ketal III will generally be prepared by acid-catalysed reaction of the ketone on which ketal III is based and not less than the equivalent amount of aliphatic C_3 - C_6 -alcohol having not less than 3 and not more than 5, preferably 3, hydroxyl groups.

25 In general, the ketal will be made by reaction of the ketone on which ketal III is based and not less than 0.5 mol equivalents of the C_3 - C_6 -alcohol having 3 to 5 hydroxyl groups will be employed. Preferably, a 1.2- to 4-fold, preferably 1.5- to 3-fold excess of this amount of the alcohol is used. Examples of acid catalysts

30 which can be used are p-toluenesulphonic acid, phosphoric acid or potassium hydrogen sulphate in catalytically effective amounts (for example 0.1 to 3 g of p-toluenesulphonic acid per mole of ketone). The reaction will preferably be carried out either in an organic solvent which together with water forms an azeotrope, so

35 that the water, which is liberated during the formation of the

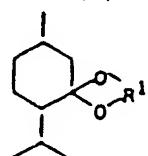
- 11 -

ketal, can be eliminated by azeotropic entrainment, or water-consuming coreagents such as, for example, trialkyl ortho esters. Examples of preferred organic solvents comprise benzene, toluene, xylene, chloroform, methylene chloride and trichloroethylene.

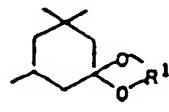
5 The reaction can be regarded as complete when water no longer separates out or when an ester/alcohol mixture is no longer separated out. It is recommended to wash the products subsequently with dilute alkali and with water, to separate and dry the
10 organic phase, to strip off the solvent and, if appropriate, to purify the residue, for example by distillation.

Particularly preferred ketals III are those of the formulae

15

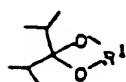


(IV)

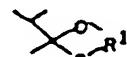


(V)

20

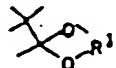


(VI)



(VII)

25



(VIII)

in which R¹ has the above-mentioned meaning, particularly preferred substances from among the ketals IV to VIII being in
30 each case the glycerol ketals.

The ketals III can have asymmetric carbon atoms; optical isomerism can therefore occur. Depending on the starting material and the preparation methods used, they can exist in the form a mixtures of the optical isomers or in the form of pure isomers.

- 12 -

The cooling effect of the isomers may differ, so that one or the other isomer may be preferred.

Aqueous Vehicle

The coolant compositions hereof, suitable for topical application to skin, will comprise coolant component I together with coolant component II in an aqueous carrier. Lower alkyl chain alcohols are common vehicle ingredients in skin care compositions. However these alcohols can cause excessive levels of a stinging or burning sensation to the skin, especially cut or irritated skin. Therefore, the compositions hereof will contain no more than about 5 30%, by weight, of C₁-C₆ monohydric alcohols, preferably no more than about 25%, more preferably no more than about 20%, even more preferably no more than about 10%, most preferably from 0% to about 15% 5%. It is especially preferred to limit the presence of C₁-C₄ monohydric alcohols, such as methanol, ethanol, propanol, isopropanol, and butanol, according to the preferred above levels.

Despite the restrictions on the use of alcohol in the present invention, the compositions hereof can provide coolant benefits upon application to the skin characterized by a strong initial 10 cooling sensation and long lasting cooling with little or no burning sensation to the skin.

The essential component of the aqueous vehicle is water. Water will generally be present at levels of at least about 50% by weight of the composition, preferably from about 75% to about 25 99.8%, more preferably from about 80% to about 97%.

The coolant components can be present at any level effective for providing coolant effect upon application to the skin. In general, the weight ratio of coolant component I to coolant component II is from about 5:1 to about 1:5, preferably from about 30 3:1 to about 1:3, more preferably from about 3:1 to about 1:1.

Coolants I and II are each typically used at levels of at least about 0.01%, by weight of the composition, preferably at least about 0.1%. The upper limit that can be used for purposes of this invention will depend upon the particular coolants being used, as excessively high levels can result in a burning sensation 35

- 13 -

upon application to the skin. In general, however, it is preferred to use coolants I and II each at levels from about 0.1% to about 1%, by weight, more preferably from about 0.1% to about 0.5%.

5 Coolant III, the optional ketal coolant, is generally used at levels of from about 0.1% to about 1.5%, more preferably from about 0.1% to about 0.75%, by weight of the composition.

Perfume

10 The compositions hereof will generally contain a perfume to impart a desired aroma, or to mask odors that may be associated with other components of the compositions. Any perfume suitable for application to the skin can be used. A wide variety of perfumes are known to those skilled in the art and are commercially available. The particular perfume used is largely a matter of choice. However the perfume should be used at a level effective for providing a noticeable aroma to the composition, or for masking undesired aroma of the composition. Also, the perfume and whatever carriers accompany it should not impart excessive stinging to the skin, especially broken or irritated skin, at the 15 levels added. In general, the compositions will comprise from about 0.1% to about 10% of a perfume component, preferably from about 0.1% to about 7%, more preferably from about 0.1% to about 3%. As used herein, perfume means aromatically active ingredients whereas perfume component includes the perfume and any 20 accompanying perfume solvent.

25 Perfumes are made by those skilled in the art in a wide variety of fragrances and strengths. Typical perfumes are described in Arctander, *Perfume and Flavour Chemicals* (Aroma Chemicals), Vol. I and II (1969); and Arctander, *Perfume and Flavour Materials of Natural Origin* (1960).

30 The perfumes selected for use herein are chosen not only for their scent and strength, but also to meet aesthetic demands of the consumer.

As disclosed in U.S. Patent 4,322,308, Hooper et al., issued 35 March 30, 1982, and U.S. Patent 4,304,679, Hooper et al., issued

- 14 -

December 8, 1981, both incorporated herein by reference, perfume components generally include, but are not limited to, volatile phenolic substances (such as iso-amyl salicylate, benzyl salicylate, and thyme oil red); essence oils (such as geranium oil, 5 patchouli oil, and petitgrain oil); citrus oils; extracts and resins (such as benzoin siam resinoid and opopanax resinoid); "synthetic" oils (such as Bergamot 37 and 430, Geranium 76 and Pomeransol 314); aldehydes and ketones (such as β -methyl naphthyl ketone, p-t-butyl-A-methyl hydrocinnamic aldehyde and p-t-amyl cyclohexanone); polycyclic compounds (such as Coumarin and 8-naphthyl methyl ether); esters (such as diethyl phthalate, phenyl-10 ethyl phenylacetate, non-anolide-1:4). Perfumes also include esters and essential oils derived from floral materials and fruits, citrus oils, absolutes, aldehydes, resinoides, musk and other animal notes (e.g., natural isolates of civet, castoreum and musk), balsamic, etc. and alcohols (such as dimyrcetol, phenylethyl alcohol and tetrahydromuguol). Examples of such components useful in perfumes herein include decyl aldehyde, undecyl aldehyde, undecylenic aldehyde, lauric aldehyde, amyl cinnamic aldehyde, ethyl methyl phenyl glycidate, methyl nonyl acetaldehyde, myristic aldehyde, nonalactone, nonyl aldehyde, octyl aldehyde, undecalactone, hexyl cinnamic aldehyde, benzaldehyde, vanillin, heliotropine, camphor para-hydroxy phenolbutanone, 6-acetyl 1,1,3,4,4,6 hexamethyl tetrahydronaphthalene, alpha-methyl ionone, 20 gamma-methyl ionone, and amyl-cyclohexanone and mixtures of these 25 components.

Perfume solvents are well known in the art, and the conventional ones can be used herein, e.g., dipropylene glycol, diethylene glycol, C₁-C₆ alcohols, etc.

30 Other Cosmetically Active Ingredients

In addition to or in place of perfume, the compositions hereof can also comprise other cosmetically active ingredients or medicaments. Cosmetically active ingredients are compounds or materials which directly affect the appearance, feel, smell, or

- 15 -

comfort of the skin, or which protect the skin from environmental factors (e.g., sun light). Medicaments are compounds or materials that have a direct medicinal or neurological effect (excluding C₁-C₆ alcohols). Such materials are well known and recognized in the art.

Cosmetically active ingredients include, but are not limited to, perfumes (as described above), skin conditioners such as skin moisturizers, humectants, and emollients, sunscreens, and pigments intended to color skin or aid in tanning of the skin. Medicaments include, but are not limited to, anti-acne ingredients, antibiotics, antimicrobials, antifungals, antivirals, antibacterials, antiprotozoals, anti-inflammatory actives, astringents, antiseptics, etc. Various preferred cosmetically active ingredients and medicaments are described in detail below.

15 Skin Conditioners

The compositions can contain one or more skin conditioners. These materials are typically used at a level of from about 0.1% to about 20%, more preferably from about 1% to about 10% and most preferably from about 2% to about 5%. Skin conditioners include, but are not limited to, moisturizers, humectants, and emollients. Exemplary moisturizers and emollients include urea; guanidine; glycolic acid and glycolate salts (e.g., ammonium and quaternary alkyl ammonium); lactic acid and lactate salts (e.g., ammonium and quaternary alkyl ammonium); aloe vera in any of its variety of forms (e.g., aloe vera gel); polyhydroxy alcohols such as sorbitol, glycerol, hexanetriol, propylene glycol, hexylene glycol and the like; polyethylene glycol; sugars and starches; sugar and starch derivatives (e.g., alkoxylated glucose); hyaluronic acid; lactamide monoethanolamine; acetamide monoethanolamine; and mixtures thereof.

Examples of suitable emollients include, but are not limited to, volatile and non-volatile silicone oils, highly branched hydrocarbons, and non-polar carboxylic acid and alcohol esters, and mixtures thereof. Emollients useful in the instant invention are further described in U.S. Patent 4,919,934, Deckner et al.,

- 16 -

issued April 24, 1990, which is incorporated herein by reference in its entirety.

The emollients, in particular, when used typically comprise in total from about 0.05% to about 50%, preferably from about 0.5% to about 25%, and more preferably from about 1% to about 10% by weight of the compositions useful in the present invention.

Preferred emollients are nonvolatile, insoluble silicone conditioning agents. The nonvolatile, insoluble silicone fluid will preferably have average viscosity of at least about 1,000, preferably from about 1,000 to about 2,000,000, centistokes at 5 25°C, more preferably from about 10,000 to about 1,800,000 centistokes, even more preferably from about 100,000 to about 1,500,000 centistokes. Lower viscosity nonvolatile silicone conditioning agents, however, can also be used, as can volatile silicones, and 10 water soluble silicones.

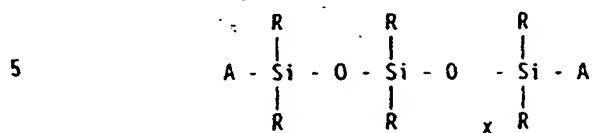
Suitable nonvolatile silicone fluids for use in hair conditioning agents include polyalkyl siloxanes, polyaryl siloxanes, polyalkylaryl siloxanes, polyether siloxane copolymer and mixtures thereof. However, other silicone fluids having hair conditioning 15 properties may be used. The nonvolatile polyalkyl siloxane fluids that may be used include, for example, polydimethylsiloxanes. These siloxanes are available, for example, from the General Electric Company as a Viscasil series and from Dow Corning as the Dow Corning 200 series. Preferably, the viscosity ranges from 20 about 10 centistokes to about 100,000 centistokes at 25°C.

The polyalkylaryl siloxane fluids that may be used, also include, for example, polymethylphenylsiloxanes. These siloxanes are available, for example, from the General Electric Company as SF 1075 methyl phenyl fluid or from Dow Corning as 556 Cosmetic Grade Fluid.

The polyether siloxane copolymer that may be used includes, for example, a polypropylene oxide modified dimethylpolysiloxane (e.g., Dow Corning DC-1248) although ethylene oxide or mixtures of 30 ethylene oxide and propylene oxide may also be used. The ethylene oxide and polypropylene oxide level must be sufficiently low to prevent solubility in water and the composition hereof.

- 17 -

Silicone fluids hereof also include polyalkyl or polaryl siloxanes with the following structure:



wherein R is alkyl or aryl, and x is an integer from about 7 to about 8,000 may be used. "A" represents groups which block the ends of the silicone chains.

The alkyl or aryl groups substituted on the siloxane chain (R) or at the ends of the siloxane chains (A) may have any structure as long as the resulting silicones remain fluid at room temperature, are hydrophobic, are neither irritating, toxic nor otherwise harmful when applied to the skin, are compatible with the other components of the composition, are chemically stable under normal use and storage conditions, and are capable of being deposited on and of conditioning skin.

Suitable A groups include methyl, methoxy, ethoxy, propoxy, and aryloxy. The two R groups on the silicone atom may represent the same group or different groups. Preferably, the two R groups represent the same group. Suitable R groups include methyl, ethyl, propyl, phenyl, methylphenyl and phenylmethyl. The preferred silicones are polydimethyl siloxane, polydiethylsiloxane, and polymethylphenylsiloxane. Polydimethylsiloxane is especially preferred.

References disclosing suitable silicone fluids include U.S. Patent 2,826,551, Geen; U.S. Patent 3,964,500, Drakoff, issued June 22, 1976; U.S. Patent 4,364,837, Pader; and British Patent 849,433, Woolston. All of these patents are incorporated herein by reference. Also incorporated herein by reference is Silicon Compounds distributed by Petrarch Systems, Inc., 1984. This reference provides an extensive (though not exclusive) listing of suitable silicone fluids.

- 18 -

Another specific silicone fluid material that can be especially useful in the silicone conditioning agents is insoluble silicone gum. The term "silicone gum", as used herein, means polyorganosiloxane materials having a viscosity at 25°C of greater than or equal to 1,000,000 centistokes. Silicone gums are described by Petrarch and others including U.S. Patent 4,152,416, Spitzer et al., issued May 1, 1979, and Noll, Walter, *Chemistry and Technology of Silicones*, New York: Academic Press 1968. Also describing silicone gums are General Electric Silicone Rubber Product Data Sheets SE 30, SE 33, SE 54 and SE 76. All of these described references are incorporated herein by reference. The "silicone gums" will typically have a mass molecular weight in excess of about 200,000, generally between about 200,000 and about 1,000,000. Specific examples include polydimethylsiloxane, (polydimethylsiloxane) (methylvinylsiloxane) copolymer, poly(dimethylsiloxane) (diphenyl siloxane)(methylvinylsiloxane) copolymer and mixtures thereof.

Other skin conditioners include oily materials, such as hydrocarbon oils and long chain esters. Suitable esters hereof include those having at least 10 carbon atoms, e.g., fatty esters, and include esters with hydrocarbyl chains derived from fatty acids or alcohols, e.g., mono-esters, polyhydric alcohol esters, and di- and tri-carboxylic acid esters. The hydrocarbyl radicals of the fatty esters hereof can also include or have covalently bonded thereto other compatible functionalities, such as amides and alkoxy moieties (e.g., ethoxy or ether linkages, etc.).

Hydrocarbon oils include cyclic hydrocarbons, straight chain aliphatic hydrocarbons (saturated or unsaturated), and branched chain aliphatic hydrocarbons (saturated or unsaturated). Straight chain hydrocarbon oils will preferably contain from about 12 to about 19 carbon atoms. Branched chain hydrocarbon oils can and typically may contain higher numbers of carbon atoms. Specific examples include paraffin oil, mineral oil, saturated and unsaturated dodecane, saturated and unsaturated tridecane, saturated and unsaturated tetradecane, saturated and unsaturated pentadecane, saturated and unsaturated hexadecane, and mixtures there-

- 19 -

of. Branched-chain isomers of these compounds, as well as of higher chain length hydrocarbons, can also be used. Exemplary branched-chain isomers are highly branched saturated or unsaturated alkanes, such as the permethyl-substituted isomers, e.g., the permethyl-substituted isomers of hexadecane and eicosane, such as 2, 2, 4, 4, 6, 6, 8, 8-dimethyl-10-methylundecane and 2, 2, 4, 4, 6, 6-dimethyl-8-methylnonane, sold by Permethyl Corporation.

5 Monocarboxylic acid esters hereof include esters of alcohols and/or acids of the formula R'COOR wherein alkyl or alkenyl radicals and the sum of carbon atoms in R' and R is at least 10, preferably at least 20.

10 Fatty esters include, for example, alkyl and alkenyl esters of fatty acids having aliphatic chains with from about 10 to about 22 carbon atoms, and alkyl and alkenyl fatty alcohol carboxylic acid esters having an alkyl and/or alkenyl alcohol-derived aliphatic chain with about 10 to about 22 carbon atoms, and combinations thereof.

15 The mono-carboxylic acid ester need not necessarily contain at least one chain with at least 10 carbon atoms, so long as the total number of aliphatic chain carbon atoms is at least 10. Examples include diisopropyl adipate, diisohexyl adipate, and diisopropyl sebacate.

20 Di- and tri-alkyl and alkenyl esters of carboxylic acids can also be used. These include, for example, esters of C₄-C₈ dicarboxylic acids such as C₁-C₂₂ esters (preferably C₁-C₆) of succinic acid, glutaric acid, adipic acid, hexanoic acid, heptanoic acid, and octanoic acid.

25 Polyhydric alcohol esters include alkylene glycol esters, for example ethylene glycol mono and di-fatty acid esters, diethylene glycol mono- and di-fatty acid esters, polyethylene glycol mono- and di-fatty acid esters, propylene glycol mono- and di-fatty acid esters, polypropylene glycol monostearate, polypropylene glycol 2000 monostearate, ethoxylated propylene glycol monostearate, glyceryl mono- and di-fatty acid esters, polyglycerol poly-fatty acid esters, ethoxylated glyceryl monostearate, 1,3-butylene glycol

- 20 -

monostearate, 1,3-butylene glycol distearate, polyoxyethylene polyol fatty acid ester, sorbitan fatty acid esters, and polyoxyethylene sorbitan fatty acid esters.

Glycerides include mono-, di-, and tri-glycerides. More 5 specifically, included are the mono-, di-, and tri-esters of glycerol and long chain carboxylic acids, such as C₁₀-C₂₂ carboxylic acids. A variety of these types of materials can be obtained from vegetable and animal fats and oils, such as castor oil, safflower oil, cottonseed oil, corn oil, olive oil, cod liver oil, 10 almond oil, avocado oil, palm oil, sesame oil, lanolin and soybean oil. Synthetic oils include triolein and tristearin glyceryl dilaurate.

15

20

25

30

35

- 21 -

Sunscreens

The compositions useful in the methods of the present invention can also optionally comprise at least one sunscreening agent. A wide variety of one or more sunscreening agents are suitable for use in the present invention and are described in U.S. Patent 5,087,445, Haffey et al., issued February 11, 1992; U.S. Patent 5,073,372, to Turner et al., issued December 17, 1991; U.S. Patent 5,073,371, Turner et al., issued December 17, 1991; and Segarin, et al., at Chapter VIII, pages 189 et seq., of *Cosmetics Science and Technology*, all of which are incorporated herein by reference in their entirety.

Preferred among those sunscreens which are useful in the compositions of the instant invention are those selected from the group consisting of ethylhexyl p-methoxycinnamate, octocrylene, octyl salicylate, oxybenzone, and mixtures thereof.

Other useful sunscreens include the solid physical sunblocks such as titanium dioxide (e.g., micronized titanium dioxide, 0.03 microns), zinc oxide, silica, iron oxide and the like.

Still other useful sunscreens are those disclosed in U.S. Patent 4,937,370, Sabatelli, issued June 26, 1990 and U.S. Patent 4,999,186, Sabatelli et al., issued March 12, 1991; these two references are incorporated by reference herein in their entirety.

Generally, the sunscreens can comprise from about 0.5% to about 20% of the compositions useful herein. Exact amounts will vary depending upon the sunscreen chosen and the desired Sun Protection Factor (SPF). SPF is a commonly used measure of photoprotection of a sunscreen against erythema. See *Federal Register*, Vol. 43, No. 166, pp 38206-38269, August 25, 1978, which is incorporated herein by reference in its entirety.

Surfactants

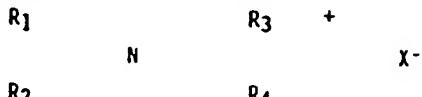
Surfactants can be used as cosmetically active ingredients for cleaning purposes, or to boost foaming properties of the composition, e.g., anionic, amphoteric, and nonionic surfactants,

- 22 -

or for skin conditioning, e.g., cationic surfactants. Anionic, amphoteric, zwitterionic and nonionic surfactants, which can also be useful as emulsifiers, are described further below.

Cationic surfactants include those that contain amino or quaternary ammonium hydrophilic moieties which are positively charged when dissolved in the aqueous composition of the present invention. Cationic surfactants among those useful herein are disclosed in the following documents, all incorporated by reference herein: M.C. Publishing Co., McCutcheon's, Detergents & Emulsifiers, (North American edition 1979); Schwartz, et al., Surface Active Agents, Their Chemistry and Technology, New York: Interscience Publishers, 1949; U.S. Patent 3,155,591, Hilfer, issued November 3, 1964; U.S. Patent 3,929,678, Laughlin, et al., issued December 30, 1975; U.S. Patent 3,959,461, Bailey, et al., issued May 25, 1976; and U.S. Patent 4,387,090, Bolich, Jr., issued June 7, 1983. If included in the compositions of the present invention, the cationic surfactant is present at from about 0.05% to about 5%.

Among the quaternary ammonium-containing cationic surfactant materials useful herein are those of the general formula:



wherein R_1-R_4 are independently an aliphatic group of from about 1 to about 22 carbon atoms, or an aromatic, alkoxy, polyoxyalkylene, alkylamido, hydroxyalkyl, aryl or alkylaryl group having from about 12 to about 22 carbon atoms; and X is an anion selected from halogen, acetate, phosphate, nitrate and alkylsulfate radicals. The aliphatic groups may contain, in addition to carbon and hydrogen atoms, ether linkages, and other groups such as amino groups. The longer chain aliphatic groups, e.g., those of about 12 carbons, or higher, can be saturated or unsaturated.

Other quaternary ammonium salts useful herein are diquaternary ammonium salts, such as tallow propane diammonium dichloride.

- 23 -

Quaternary ammonium salts include dialkyldimethyl-ammonium chlorides, wherein the alkyl groups have from about 12 to about 22 carbon atoms and are derived from long-chain fatty acids, such as hydrogenated tallow fatty acid (tallow fatty acids yield quaternary compounds wherein R₁ and R₂ have predominately from 16 to 18 carbon atoms). Di-(saturated or unsaturated tallow) dimethyl ammonium salts are particularly preferred.

Salts of primary, secondary and tertiary fatty amines are also suitable cationic surfactant materials. The alkyl groups of such amines preferably have from about 12 to about 22 carbon atoms, and may be substituted or unsubstituted. Such amines, useful herein, include stearamido propyl dimethyl amine, diethyl amino ethyl stearamide, dimethyl stearamine, dimethyl soyamine, soyamine, myristyl amine, tridecyl amine, ethyl stearylamine, N-tallowpropane diamine, ethoxylated (5 moles E.O.) stearylamine, dihydroxy ethyl stearylamine, and arachidyl behenylamine. Suitable amine salts include the halogen, acetate, phosphate, nitrate, citrate, lactate and alkyl sulfate salts. Such salts include stearylamine hydrochloride, soyamine chloride, stearylamine formate, N-tallowpropane diamine dichloride and stearamidopropyl dimethylamine citrate. Cationic amine surfactants included among those useful in the present invention are disclosed in U.S. Patent 4,275,055, Nachtigal, et al., issued June 23, 1981, incorporated by reference herein.

25 Cationic Polymer Conditioning Agent

The compositions of the present invention can also comprise a cationic organic polymer conditioning agent. The polymeric cationic hair conditioning agent hereof will generally be present at levels of from about 0.05% to about 5%, preferably from about 0.1% to about 4%, more preferably from about 0.2% to about 3%, by weight, of the shampoo composition. Preferably the cationic organic polymer is sufficiently soluble in water to form a substantially clear solution to the naked eye at a concentration of 0.1% in water (distilled or equivalent) at 25°C.

- 24 -

The cationic polymers hereof will generally have a weight average molecular weight which is at least about 5,000, typically at least about 10,000, and is less than about 10 million. Preferably, the molecular weight is from about 100,000 to about 2 million.

5 The cationic polymers typically will have cationic nitrogen-containing moieties such as quaternary ammonium or cationic amino moieties, or a mixture thereof. The precise cationic charge density is not believed to be critical to the invention. However, 10 for practical reasons, the charge density should be of a level such that efficient substantivity between the polymer and the hair can be attained. Any anionic counterions can be utilized for the cationic polymers so long as the water solubility criteria is met. Suitable counterions include halides (e.g., Cl, Br, I, or F, 15 preferably Cl, Br, or I), sulfate, and methylsulfate. Others can also be used, as this list is not exclusive. Generally, it is preferred that cationic charge density be at least about 0.2 meq/gram, more preferably at least about 0.4 meq/gram, at the pH of intended use.

20 The cationic nitrogen-containing moiety will be present generally as a substituent, on a fraction of the total monomer units of the cationic hair conditioning polymers. Thus, the cationic polymer can comprise copolymers, terpolymers, etc. of quaternary ammonium or cationic amine-substituted monomer units 25 and other non-cationic units referred to herein as spacer monomer units. Such polymers are known in the art, and a variety can be found in the *CTFA Cosmetic Ingredient Dictionary*, 3rd edition, edited by Estrin, Crosley, and Haynes, (The Cosmetic, Toiletry, and Fragrance Association, Inc., Washington, D.C., 1982).

30 Suitable cationic polymers include, for example, copolymers of vinyl monomers having cationic amine or quaternary ammonium functionalities with water soluble spacer monomers such as acrylamide, methacrylamide, alkyl and dialkyl acrylamides, alkyl and dialkyl methacrylamides, alkyl acrylate, alkyl methacrylate, vinyl 35 caprolactone, and vinyl pyrrolidone. The alkyl and dialkyl

- 25 -

substituted monomers preferably have C₁-C₃ alkyl groups, more preferably C₁-C₂ alkyl groups. Other suitable spacer monomers include vinyl esters, vinyl alcohol (made by hydrolysis of polyvinyl acetate), maleic anhydride, propylene glycol, and ethylene glycol.

The cationic amines can be primary, secondary, or tertiary amines, depending upon the particular species and the pH of the shampoo. In general, secondary and tertiary amines, especially tertiary amines, are preferred.

Suitable cationic amino and quaternary ammonium monomers include, for example, vinyl compounds substituted with dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, monoalkylaminoalkyl acrylate, monoalkylaminoalkyl methacrylate, trialkylmethacryloxyalkyl ammonium salt, trialkyl acryloxyalkyl ammonium salt, diallyl quaternary ammonium salts, and vinyl quaternary ammonium monomers having cyclic cationic nitrogen-containing rings such as pyridinium, imidazolium, and quaternized pyrrolidone, e.g., alkyl vinyl imidazolium, alkyl vinyl pyridinium, alkyl vinyl pyrrolidone salts. The alkyl portions of these monomers are preferably lower alkyls such as the C₁-C₃ alkyls, more preferably C₁ and C₂ alkyls.

Suitable amine-substituted vinyl monomers for use herein include dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, dialkylaminoalkyl acrylamide, and dialkylaminoalkyl methacrylamide, wherein the alkyl groups are preferably C₁-C₃ hydrocarbyls, more preferably C₁-C₂ alkyls.

The cationic polymers hereof can comprise mixtures of monomer units derived from amine- and/or quaternary ammonium-substituted monomer and/or compatible spacer monomers.

Suitable cationic hair conditioning polymers include, for example: copolymers of 1-vinyl-2-pyrrolidone and 1-vinyl-3-methylimidazolium salt (e.g., chloride salt) (referred to in the industry by the Cosmetic, Toiletry, and Fragrance Association, "CTFA", as Polyquaternium-16), such as those commercially available from BASF Wyandotte Corp. (Parsippany, NJ, USA) under the LUVIQUAT

- 26 -

tradename (e.g., LUVIQUAT FC 370 and FC 905); copolymers of 1-vinyl-2-pyrrolidone and dimethylaminoethyl methacrylate (referred to in the industry by CTFA as Polyquaternium-11) such as those commercially available from Gaf Corporation (Wayne, NJ, USA) under the GAFQUAT tradename (e.g., GAFQUAT 755N); cationic diallyl quaternary ammonium-containing polymers, including, for example, dimethyldiallylammonium chloride homopolymer and copolymers of acrylamide and dimethyldiallylammonium chloride, referred to in the industry (CTFA) as Polyquaternium 6 and Polyquaternium 7, respectively, such materials being available from Merck and Co., Inc. in their Merquat® series; copolymers of acrylamide and dimethyl diallyl ammonium chloride, such as those available under the MERQUAT tradename from Calgon Corp. (Pittsburgh, PA, USA) (e.g., MERQUAT 550); and mineral acid salts of amino-alkyl esters of homo- and co-polymers of unsaturated carboxylic acids having from 3 to 5 carbon atoms, as described in U.S. Patent 4,009,256, incorporated herein by reference.

Other cationic polymers that can be used include polysaccharide polymers, such as cationic cellulose derivatives and cationic starch derivatives.

Cationic polysaccharide polymer materials suitable for use herein include those of the formula:



wherein: A is an anhydroglucose residual group, such as a starch or cellulose anhydroglucose residual,

R is an alkylene oxyalkylene, polyoxalkylene, or hydroxyalkylene group, or combination thereof,

R¹, R², and R³ independently are alkyl, aryl, alkylaryl, arylalkyl, alkoxyalkyl, or alkoxyaryl groups, each group containing up to about 18 carbon atoms, and the total number of carbon atoms for each cationic moiety (i.e., the sum of

- 27 -

carbon atoms in R¹, R² and R³) preferably being about 20 or less, and

X is an anionic counterion, as previously described.

The degree of cationic substitution is typically from about 0.01-1
5 cationic groups per anhydroglucose unit.

Cationic cellulose is available from Amerchol Corp. (Edison, NJ, USA) in their Polymer JR® and LR® series of polymers, as salts of hydroxyethyl cellulose reacted with trimethyl ammonium substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 10. Another type of cationic cellulose includes the polymeric quaternary ammonium salts of hydroxyethyl cellulose reacted with lauryl dimethyl ammonium-substituted opoxide, referred to in the industry (CTFA) as Polyquaternium 24. These materials are available from Amerchol Corp. (Edison, NJ, USA) under the trade-name Polymer LM-200.
10
15

Other cationic polymers that can be used include cationic guar gum derivatives, such as guar hydroxypropyltrimonium chloride (commercially available from Celanese Corp. in their Jaguar® series). Other materials include quaternary nitrogen-containing
20 cellulose ethers (e.g., as described in U.S. Patent 3,962,418, incorporated by reference herein), and copolymers of etherified cellulose and starch (e.g., as described in U.S. Patent 3,958,581, incorporated by reference herein).

Other cosmetic actives include, but are not limited to: skin penetration aids such as DMSO, 1-dodecyl-azacycloheptan-2-one (available as Azone from the Upjohn Co.) and the like; artificial tanning agents such as dihydroxyacetone and the like; skin bleaching (or lightening) agents including but not limited to hydroquinone, ascorbic acid, kojic acid and sodium metabisulfite;
25
30
35

antioxidants; and agents which soothe skin or aid in healing of irritated skin, nonlimiting examples of these aesthetic components include panthenol, Clove oil, camphor, eucalyptus oil, eugenol, menthol lactate, witch hazel distillate, allantoin, bisabolol, and the like.

- 28 -

Medicaments

Medicaments that can be added include anti-acne ingredients such as salicylic acid, pantothenic acid and pantothenic acid derivatives (e.g., alcohol, aldehyde, alcohol ester, acid ester derivatives, etc., especially alcohol derivatives such as panthenol. A safe and effective amount of the medicament is included in the compositions to achieve the intended medical effect at the expected unit dosage. The medicaments are typically used at levels of about 0.1% to about 10% by weight of the composition.

Examples of other medicaments include keratolytics such as sulfur, lactic acid, glycolic, pyruvic acid, urea, resorcinol, and N-acetylcysteine; retinoids such as retinoic acid and its derivatives (e.g., cis and trans); antibiotics, antimicrobials, antibacterials, antifungals, antiprotozoals, and antivirals (e.g., benzoyl peroxide, octopirox, erythromycin, tetracyclin, triclosan, azelaic acid and its derivatives, phenoxy ethanol and phenoxy proponol, ethylacetate, clindamycin and mecloxycline, triclosan, chlorhexidine, tetracycline, neomycin, miconazole hydrochloride, octopirox, parachlorometaxylenol, nystatin, tolnaftate, clotrimazole, and the like); sebastats such as flavinoids; hydroxy acids; antipruritic drugs including, for example, pharmaceutically-acceptable salts of methdilizine and trimipramine; and bile salts such as scymnol sulfate and its derivatives, deoxycholate, and cholate.

Also, useful are non-steroidal anti-inflammatory drugs (NSAIDS). The NSAIDS can be selected from the following categories: propionic acid derivatives; acetic acid derivatives; fenamic acid derivatives; biphenylcarboxylic acid derivatives; and oxicams. All of these NSAIDS are fully described in the U.S. Patent 4,985,459 to Sunshine et al., issued January 15, 1991, incorporated by reference herein. Most preferred are the propionic NSAIDS including but not limited to aspirin, acetaminophen, ibuprofen, naproxen, benoxaprofen, flurbiprofen, fenoprofen, fenbufen, ketoprofen, indoprofen, pirprofen, carprofen, oxaprozin, pranoprofen, miroprofen, tioxaprofen, suprofen, alminoprofin,

- 29 -

tiaprofenic acid, fluprofen and bucloxic acid. Also useful are the steroidal anti-inflammatory drugs including hydrocortisone and the like.

Coolant Compositions

5 The compositions hereof can be used for a wide range of topical skin care compositions. They can be in the form of liquids, creams, lotions, and gels. Specific examples include toiletries such as after-shave lotions (i.e., post-shave lotions), pre-shave lotions, shaving aids such as shaving creams, gels and
10 lotions, deodorants, perfumes, colognes, liquid soap or detergents, face creams, hand creams, skin creams, sunburn lotions, and sunscreen lotions. Medicament products include first aid and antiseptic products, anti-acne ointments, and topical anti-inflammatory compositions.

15 The aqueous vehicle can contain ingredients to modify the physical properties of the composition, such as thickeners, gelling agents, and viscosity modifiers. The compositions can also contain other ingredients, such as pigments, preservatives, pH modifiers, etc. In general, it is preferred for pH of the
20 compositions to be between about pH 5.5 and about pH 8.5 in order to minimize burning or irritation of the skin although it is not intended to necessarily limit this invention to such range. Certain products, such as astringents containing acidic agents, may need to be outside this range for optional effectiveness.

25 However, such products are still intended to be encompassed herein since long lasting cooling with relatively low skin burning and irritation can still be obtained.

30 The essential coolants hereof are insoluble in water. It is therefore preferred to incorporate them into the compositions in a way so that they remain stably distributed throughout the composition. This can be done by forming a dispersion, emulsion, or microemulsion, with the coolants solubilized in a suitable solvent that then forms the dispersion or emulsion (including microemulsions) in the aqueous carrier. Emulsion systems include oil-in-water, water-in-oil, water-in-oil-in-water, and oil-in-water-in-silicone emulsions. The emulsions can cover a broad range of
35

- 30 -

consistencies including thin lotions (which can also be suitable for spray or aerosol delivery), creamy lotions, light creams, heavy creams, and the like. Suitable solvents for the coolants include lipophilic or nonpolar solvents such as diethylene glycol, dipropylene glycol, C₁-C₆ alcohols, acetone, and other organic solvents, many of which are also suitable for use as perfume solvents. Accordingly, the perfume and coolant components can be combined in a common solvent stage which is then incorporated into the composition.

Emulsion forming techniques are well known in the art. Suitable emulsifiers for use herein are anionic, nonionic, cationic, amphoteric, and zwitterionic surfactants. Surfactants can also be included herein, at suitable levels, as cosmetically active ingredients. Cationic surfactants, e.g., quaternary ammonium compounds, can be effective skin conditioners. Anionic, nonionic, amphoteric, and zwitterionic surfactants can be effective cleaning ingredients.

The compositions useful in the methods of the present invention can optionally comprise one or more surfactants. The surfactants can be present at a level from about 0.1% to about 20%, more preferably from about 0.2% to about 10%, and most preferably from about 1% to about 7.5%. Examples of a broad variety of surfactants useful herein are described in McCutcheon's, *Detergents and Emulsifiers*, North American Edition (1986), published by Allured Publishing Corporation, which is incorporated herein by reference in its entirety.

Suitable surfactants include, but are not limited to, nonionic surfactants such as polyalkylene glycol ethers of fatty alcohols. Suitable anionic surfactants include taurates, alkyl sulfates, alkyl ethoxylated sulfates, and alkyl phosphates and alkyl ethoxylated phosphates. Nonlimiting examples of these surfactants include isoceteth-20, sodium methyl cocoyl taurate, sodium methyl oleoyl taurate, and sodium lauryl sulfate. See U.S. Patent 4,800,197, to Kowcz et al., issued January 24, 1989, which is incorporated herein by reference in its entirety.

SUBSTITUTE SHEET

CA 002477

- 31 -

Suitable alkyl and alkyl ethoxylated phosphates are of the formulas $ROPO_4M$ and $RO(C_2H_4O)_xPO_4M$ wherein R is alkyl or alkenyl of from about 8-24 carbon atoms, x is 1 to 10, and M is a water soluble cation such as ammonium, or alkali or alkaline earth metal such as sodium, potassium, and magnesium, or an amine such as triethanolamine. These are also referred to as phosphate esters.

Suitable alkyl and alkyl ethoxylated sulfates are of the formulas $ROSO_3M$ and $RO(C_2H_4O)_xSO_3M$ wherein R is alkyl or alkenyl of from about 8-24 carbon atoms, x is 1 to 10, and M is a water soluble cation such as ammonium, or alkali or alkaline earth metal such as sodium, potassium, and magnesium, or an amine such as triethanolamine.

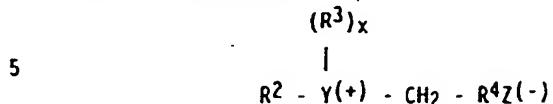
Other anionic surfactants include succinamates, e.g., disodium N-octadecylsulfosuccinamate, tetrasodium N-(1,2-dicarboxyethyl)-N-octadecylsulfosuccinamate, and dioctyl, dihexyl, and diamyl esters of sodium sulfosuccinic acid.

Suitable nonionic surfactants include: polyethylene oxide condensates of alkyl phenols, e.g., the condensation products of C₆-C₂₀ alkyl phenols with ethylene oxide, with a molar ratio of from about 10 to about 60 moles, ethylene oxide per mole of alkyl phenol; condensation products of ethylene oxide with the reaction product of propylene oxide and ethylene diamine, e.g., compounds with about 40%-80% polyoxyethylene by weight having a molecular weight of from about 5,000-11,000, wherein the ethylene diamine/propylene oxide portion has a molecular weight of about 2,500-3,000; condensation products of C₈-C₁₈ aliphatic alcohols (straight or branched chain) with ethylene oxide, with, for example about 10-30 moles of ethylene oxide per mole of alcohol; polysorbates; alkylpolyglycosides; polyethylene glycol glyceryl fatty esters; etc.

Zwitterionic surfactants are exemplified by those which can be broadly described as derivatives of aliphatic quaternary ammonium, phosphonium, and sulfonium compounds, in which the aliphatic radicals can be straight or branched chain, and wherein one of the aliphatic substituents contains from about 8 to about

- 32 -

18 carbon atoms and one contains an anionic group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate. A general formula for these compounds is:



wherein R² contains an alkyl, alkenyl, or hydroxy alkyl radical of from about 8 to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties and from 0 to about 1 glyceryl moiety; Y is selected from the group consisting of nitrogen, phosphorus, and sulfur atoms; R³ is an alkyl or monohydroxyalkyl group containing about 1 to about 3 carbon atoms; X is 1 when Y is a sulfur atom, and 2 when Y is a nitrogen or phosphorus atom; R⁴ is an alkylene or hydroxyalkylene of from about 1 to about 4 carbon atoms and Z is a radical selected from the group consisting of carboxylate, sulfonate, sulfate, phosphonate, and phosphate groups.

10

Examples of amphoteric surfactants which can be used in the compositions of the present invention are those which are broadly described as derivatives of aliphatic secondary and tertiary amines in which the aliphatic radical can be straight or branched chain and wherein one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic water solubilizing group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate. Examples of compounds falling within this definition are sodium 3-dodecyl-aminopropionate, sodium 3-dodecylamino-propane sulfonate, sodium lauryl sarcosinate, N-higher alkyl aspartic acids such as those produced according to the teaching of U.S. Patent 2,438,091, and the products sold under the trade name "Miranol" and described in U.S. Patent 2,528,378.

15

Other amphoteric surfactants also include sultaines and amidosultaines. Sultaines, including amidosultaines, include for example, cocodimethylpropylsultaine, stearyltrimethylpropylsultaine, lauryl-bis-(2-hydroxyethyl) propylsultaine and the like; and the amidosultaines such as cocoamidodimethylpropylsultaine, stearyltrimidodimethylpropylsultaine, lauryltrimidobis-(2-hydroxy-

20

25

30

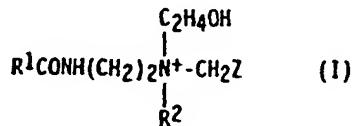
35

- 33 -

ethyl) propylsultaine, and the like. Preferred are amidohydroxysultaines such as the C₁₂-C₁₈ hydrocarbyl amidopropyl hydroxysultaines, especially C₁₂-C₁₄ hydrocarbyl amido propyl hydroxysultaines, e.g., laurylamidopropyl hydroxysultaine and cocamido-propyl hydroxysultaine. Other sultaines are disclosed in U.S. Patent 3,950,417, issued April 13, 1976, incorporated herein by reference.

Examples of betaines useful herein include the high alkyl betaines, such as coco dimethyl carboxymethyl betaine, cocoamido-propyl betaine, cocobetaine, lauryl amidopropyl betaine, oleyl betaine, lauryl dimethyl carboxymethyl betaine, lauryl dimethyl alphacarboxyethyl betaine, cetyl dimethyl carboxymethyl betaine, lauryl bis-(2-hydroxyethyl) carboxymethyl betaine, stearyl bis-(2-hydroxypropyl) carboxymethyl betaine, oleyl dimethyl gamma-carboxypropyl betaine, and lauryl bis-(2-hydroxypropyl)alpha-carboxyethyl betaine. The sulfobetaines may be represented by coco dimethyl sulfopropyl betaine, stearyl dimethyl sulfopropyl betaine, lauryl dimethyl sulfoethyl betaine, lauryl bis-(2-hydroxyethyl) sulfopropyl betaine and the like; amidobetaines and amidosulfobetaines, wherein the RCONH(CH₂)₃ radical is attached to the nitrogen atom of the betaine are also useful in this invention. Preferred betaines for use in the present compositions are cocoamidopropyl betaine, cocobetaine, lauryl amidopropyl betaine, and oleyl betaine.

Other specific amphoteric materials include imidazolinium materials depicted by formula:



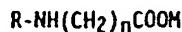
wherein R¹ is C₈-C₂₂ alkyl or alkenyl, R² is hydrogen, CO₂M, CH₂CO₂M, or CH₂CH₂M, Z is CO₂M or CH₂CO₂M, and M is hydrogen, alkali metal, alkaline earth metal, ammonium or alkanol ammonium.

Materials included are cocoamphocarboxypropionate, cocoamphocarboxy propionic acid, and cocoamphocarboxyglycinate. Mixtures

- 34 -

of these materials may also be used. The most preferred material of this type for use in the present invention is cocoamphocarboxy-glycinate (also known as cocoamphodiacetate).

Another specific class of amphoteric surfactants is defined
5 by the aminoalkanoates of formula:



and the iminodialkanoates of formula:



and mixtures thereof; wherein n and m are numbers from 1 to 4, R is C₈-C₂₂ alkyl or alkenyl, and M is hydrogen, alkali metal, alkaline earth metal, ammonium or alkanolammonium.

15 Examples of such amphoteric surfactants include n-alkyl-amino-propionates, n-alkyliminodipropionates, and mixtures thereof.

Gelling Agents and Thickeners

20 Optional components of the compositions useful in the instant invention also include gelling agents and thickeners. A preferred gelling agent is a carboxylic copolymer (acrylic acid copolymer), for example Carbomer 954 and Carbomer 1342 (available as Carbopol 954 and Carbopol 1342 from B. F. Goodrich). These polymers are more fully described in U.S. Patent 4,509,949, to Huang et al., issued April 5, 1985, and U.S. Patent 2,798,053, to Brown, issued July 2, 1957, both of which are incorporated herein by reference in their entirety. Also useful are the acrylate/alkyl acrylate crosspolymers such as Acrylates/C10-C30 Alkyl Acrylate Cross-
25 polymer (available as Pemulen TR-1 and Pemulen TR-2 from Goodrich).

30 These polymers can be used at levels from about 0.015% to about 0.75%, preferably from about 0.05% to about 0.25% and most preferably from about 0.075% to about 0.175% in the compositions herein.

- 35 -

Other gellants and thickeners include alkyl glycols, alkyl modified cellulose polymers, gums such as guar gums and xanthan gums, and mixtures thereof. Exemplary alkyl modified cellulose polymers, specifically those selected from the group consisting of 5 methylcellulose, ethylcellulose, hydroxybutyl methylcellulose, hydroxy ethylcellulose, hydroxy propylcellulose, hydroxypropyl methylcellulose cellulose, and mixtures thereof.

Suitable thickening systems are also disclosed in U.S. Patent 5,106,609, Bolich et al., issued April 21, 1992, U.S. Patent 10 5,104,646, Bolich et al., issued April 14, 1992, U.S. Patent 5,100,658, Bolich et al., issued March 31, 1992, and U.S. Patent 5,100,657, Ansher-Jackson et al., issued March 31, 1992, all of which are incorporated herein by reference. These patents disclose thickened vehicle systems containing water, a surfactant 15 or water insoluble polymer, and a nonionic, hydrophobically modified water soluble polymer, such as a nonionic, long chain alkylated (e.g., hydroxyalkyl, urethane, or acyl radicals) cellulose polymer, e.g., Natrosol Plus CS Grade 67, a hydrophobically modified (cetylated) hydroxyethyl cellulose available from 20 Aqualon Corporation, Wilmington, Delaware, USA. Such compositions typically comprise from about 0.1% to about 10% of the polymer, from 0.02% to about 10.0% of a surfactant (preferably a water-insoluble surfactant, and also preferably no more than about 2% water soluble surfactant) or from about 0.3% to about 5% of a 25 water soluble polymeric thickener (e.g., locust bean and guar gums), and water.

Other aqueous vehicle systems are gels based upon water, a lipid or lipid-type ingredient (e.g., fatty alcohol), and a cationic surfactant.

30 Suspending agents also include long chain acyl derivative materials, long chain amine oxides, or mixtures of such materials. Preferably such suspending agents are present in the composition in crystalline form. Suspending agents of this type are described in U.S. Patent 4,741,855, Grote and Russell, issued May 3, 1988, 35 incorporated herein by reference. Included are ethylene glycol

- 36 -

esters of fatty acids preferably having from about 16 to about 22 carbon atoms. More preferred are the ethylene glycol stearates, both mono and distearate, but particularly the distearate. Other suspending agents found useful are alkanol amides of fatty acids, 5 preferably having from about 16 to about 22 carbon atoms, more preferably about 16 to 18 carbon atoms. Other long chain acyl derivatives include long chain esters of long chain fatty acids (e.g., stearyl stearate, cetyl palmitate, etc.); glyceryl esters (e.g., glyceryl distearate) and long chain esters of long chain 10 alkanol amides (e.g., stearamide diethanolamide distearate, stearamide monoethanolamide stearate). Long chain acyl derivatives also include N,N-dihydrocarbyl amido benzoic acid and soluble salts thereof (e.g., Na and K salts), particularly N,N-di(hydrogenated) C₁₆, C₁₈ and tallow amido benzoic acid species of 15 this family, which are commercially available from Stepan Company (Northfield, Illinois, USA). Long chain acyl derivatives, ethylene glycol esters of long chain carboxylic acids, long chain amine oxides, and alkanol amides of long chain carboxylic acids in addition to the preferred materials listed above may be used as 20 suspending agents. For example, it is contemplated that suspending agents with long chain hydrocarbys having C₈-C₂₂ chains may be used.

Suspending agents also include long chain amine oxides such 25 as alkyl (C₁₆-C₂₂) dimethyl amine oxides, e.g., stearyl dimethyl amine oxide.

The acyl derivative and amine oxide suspending agents are typically present in pourable, liquid formulations at a level of from about 0.1% to about 5.0%, preferably from about 0.5% to about 3.0%.

30 Another type of suspending agent that can be used is xanthan gum. Shampoo compositions utilizing xanthan gum as a suspending agent for the silicone hair conditioning component are described in U.S. Patent 4,788,006, Bolich and Williams, issued November 29, 1988, incorporated herein by reference. Xanthan gum is bio- 35 synthetic gum material that is commercially available. It is a

- 37 -

heteropolysaccharide with a molecular weight of greater than 1 million. It is believed to contain D-glucose, D-mannose and D-glucuronate in the molar ratio of 2.8:2.0:2.0. The polysaccharide is partially acetylated with 4.7% acetyl. This information and other is found in Whistler, Roy L. Editor *Industrial Gums - Polysaccharides and Their Derivatives* New York: Academic Press, 1973. Kelco, a Division of Merck & Co., Inc. offers xanthan gum as Keltrol®. The gum, when used as the silicone hair conditioning component suspending agent, will typically be present in pourable, liquid formulations at a level of from about 0.3% to about 3%, preferably from about 0.4% to about 1.2% in the compositions of the present invention.

Combinations of long chain acyl derivatives and xanthan gum are disclosed as a suspending agent for silicone hair conditioners in U.S. Patent 4,704,272, Oh et al., issued November 3, 1987, incorporated herein by reference, and may also be used in the present compositions.

Another type of suspending agent that can be used is carboxyvinyl polymer. Preferred polymers are copolymers of acrylic acid crosslinked with polyallylsucrose as described in U.S. Patent 2,798,053, Brown, issued July 2, 1957, incorporated herein by reference. These polymers are provided by B. F. Goodrich Company as, for example, Carbopol 934, 940, 941, and 956.

A carboxyvinyl polymer is an interpolymer of a monomeric mixture comprising a monomeric olefinically unsaturated carboxylic acid, and from about 0.1% to about 10% by weight of the total monomers of a polyether of a polyhydric alcohol, which polyhydric alcohol contains at least four carbon atoms to which are attached at least three hydroxyl groups, the polyether containing more than one alkenyl group per molecule. Other monoolefinic monomeric materials may be present in the monomeric mixture if desired, even in predominant proportion.

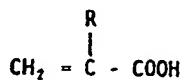
Preferred polyhydric alcohols used to product carboxyvinyl polymers include polyols selected from the class consisting of oligosaccharides, reduced derivatives thereof in which the

- 38 -

carbonyl group is converted to an alcohol group, and pentaerythritol; more preferred are oligosaccharides, most preferred is sucrose. It is preferred that the hydroxyl groups of the polyol which are modified be etherified with allyl groups, the polyol having at least two allyl ether groups per polyol molecule.

5 When the polyol is sucrose, it is preferred that the sucrose have at least about five allyl ether groups per sucrose molecule. It is preferred that the polyether of the polyol comprise from about 0.1% to about 4% of the total monomers, more preferably from about 10 0.2% to about 2.5%.

Preferred monomeric olefinically unsaturated carboxylic acids for use in producing carboxyvinyl polymers used herein include monomeric, polymerizable, alpha-beta monoolefinically unsaturated lower aliphatic carboxylic acids; more preferred are monomeric 15 monoolefinic acrylic acids of the structure



where R is a substituent selected from the group consisting of 20 hydrogen and lower alkyl groups; most preferred is acrylic acid.

Preferred carboxyvinyl polymers used in formulations of the present invention have a molecular weight of at least about 750,000; more preferred are carboxyvinyl polymers having a molecular weight of at least about 1,250,000; most preferred are carboxyvinyl polymers having a molecular weight of at least about 25 3,000,000.

Materials that can also be used as suspension or gelling agents, include water soluble or colloidally water soluble polymers like cellulose ethers (e.g., hydroxyethyl cellulose), guar gum, polyvinyl alcohol, polyvinyl pyrrolidone, hydroxypropyl guar 30 gum, starch and starch derivatives. In general, these are used at a level of from about 0.1% to about 10%, most commonly from about 0.3% to about 5.0% by weight of the composition.

Other Optional Components

35 A variety of additional ingredients can be incorporated into the compositions useful in the methods of the present invention.

- 39 -

Non-limiting examples of these additional ingredients include polymers for aiding the film-forming properties and substantivity of the composition (such as a copolymer of eicosene and vinyl pyrrolidone, an example of which is available from GAF Chemical Corporation as Ganex V-220®); preservatives; colors and dyes; and sequestrants and chelators.

The compositions hereof can be dispensed from conventional containers or bottles, aerosol or nonaerosol spray containers, tubes, etc. Additionally, the compositions herein can be incorporated into a pad, which is used by rubbing or wiping against the skin. Preferably these pads comprise from about 50% to about 75% by weight of one or more layers of nonwoven fabric material and from about 20% to about 75% by weight (on dry solids basis) of a water soluble polymeric resin. Such pads are described in detail in U.S. Patent 4,891,228, to Thaman et al., issued January 2, 1990 and U.S. Patent 4,891,227, to Thaman et al., issued January 2, 1990; both of which are incorporated by reference herein.

The present invention further encompasses a method for providing a cooling sensation to the skin comprising applying an effective amount of any of the compositions hereof to the skin. The composition is left on the skin to provide an effective, long-lasting cooling sensation. Typically, from about 0.05 g to about 10 g per cm² of skin is applied.

EXAMPLES

The following examples serve to further describe and demonstrate embodiments within the scope of the invention, but are not to be construed as limitations of the present invention as many variations are possible without departing from the spirit of the invention. The scope of the invention is defined in the claims which follow.

Examples 1-5

Given below are examples of after shave lotions of the present invention.

- 40 -

	<u>Ingredients (wt. %)</u>	<u>Examples</u>				
		1	2	3	4	5
	Ethanol	20.00	0.00	10.00	30.00	5.00
	Perfume Component	3.00	1.50	2.00	5.00	1.00
5	Phosphate Ester 1	7.50	6.00	5.00	7.50	2.00
	Sorbitan Monooleate	5.00	2.00	2.50	4.00	1.00
	Glydant Plus 2	0.15	0.20	0.18	0.10	0.18
	Disodium EDTA 3	0.10	0.10	0.10	0.10	0.10
	Butylene Glycol	5.00	1.00	3.00	5.00	1.00
10	Coolant 1 4	0.08	0.16	0.14	0.12	0.06
	Coolant 2 5	0.08	0.08	0.06	0.04	0.12
	Water	- - - - q.s. to 100% - - - -				

1 Potassium Butylcarbamol Phosphate

2 DMDM hydantoin and iodopropynyl butylcarbamate, available from Lonza, Inc. (Fairlawn, NJ, USA)

3 Disodium ethylenediaminetetraacetate

4 N,2,3-trimethyl-2-isopropylbutanamide

5 N-ethyl p-menthan-3-carboxamide

The above compositions are made by mixing Coolants 1 and 2, the perfume component, and phosphate ester surfactant to form a solution. A second solution is formed by mixing the disodium EDTA, Glydant Plus, butylene glycol, ethanol, and water. The first and second mixtures are then combined and mixed to form the final product.

The compositions can provide an effective, long-lasting cooling sensation to the skin with low or zero noticeable stinging to the skin, including cut or irritated skin.

Examples 6-10

Given below are examples of astringent cleaning compositions of the present invention.

	<u>Ingredients (wt. %)</u>	<u>Examples</u>				
		6	7	8	9	10
	Ethanol	30.00	30.00	20.00	30.00	20.00
	Salicylic Acid	0.35	0.40	0.30	0.50	0.35
35	Perfume Component	0.10	0.10	0.05	0.08	0.15

- 41 -

PEG (40) Hydrogenated

	Castor Oil 1	0.03	0.05	0.15	0.10	0.20
	Coolant 1 2	0.08	0.12	0.12	0.05	0.12
	Coolant 2 3	0.04	0.04	0.08	0.05	0.12
5	Coolant 3 4 (Optional)	0.12	0.08	0.12	0.25	0.12
	Water	- - - - -	q.s to 100%	- - - - -		

1 Polyethylene (40) glycol hydrogenated castor oil

2 N,1,3-trimethyl-2-isopropylbutanamide

3 N-ethyl p-menthan-3-carboxamide

4 1-Menthon-/d-iso-menthon glycerin ketal

10 The perfume component, salicylic acid, ethanol, coolants, and surfactant are combined and mixed together to form a solution. This is then added to the water and mixed to form the final product. Optionally, Coolant 3 can be included in the compositions.

15 The compositions can provide an effective, long lasting cooling sensation to the skin, along with astringent benefits, with relatively low skin sting.

Example 11

20 Given below is a shave cream composition of the present invention.

	Ingredients	<u>Example 11</u>
	Stearic Acid	6.30
	Lauramide DEA 1	1.00
25	Triethanolamine	3.90
	Perfume Component	0.25
	Isobutane	2.80
	Propane	0.50
	Coolant 1	0.16
30	Coolant 2	0.12
	Water	q.s. to 100 %

1 Lauric diethanolamide

35 The shave cream is prepared by mixing and heating the water, stearic acid, lauramide DEA, and triethanolamine to about 75°C to form a solution. The mixture is cooled to room temperature. The

- 42 -

perfume component and sensates are then mixed together and added to the cooled mixture. This mixture is then dispensed in a conventional shave cream aerosol container along with the isobutane and propane propellants. Coolants 1 and 2 are as defined above. Optionally, Coolant 3 (as defined above) can also be added, e.g., at a level of about 0.12%.

The composition hereof can provide excellent shaving performance upon application to the face and shaving, along with providing a long-lasting cooling sensation to the skin, with low skin sting.

10

Example 12

Given below is a shave gel composition of the present invention.

	<u>Ingredients (wt. %)</u>	<u>Example 12</u>
15	Palmitic Acid	7.20
	Triethanolamine	5.30
	Stearic Acid	2.40
	Isopentane	2.00
	PEG-14M 1	0.25
20	Isobutane	0.35
	Perfume Component	0.25
	Coolant 1	0.16
	Coolant 2	0.06
	Water	q.s. to 100%

25 1 Polyethylene glycol (20,000 d.p.)

The shave gel is prepared by mixing and heating the water, palmitic acid, stearic acid, triethanolamide, and PEG-14M to about 75°C to form a solution. The mixture is cooled to room temperature. The perfume component and coolants are mixed together to form a solution, which is then added to and mixed with the first mixture. Coolants 1 and 2 are as previously defined. Optionally, 0.12% of Coolant 3 can also be included. The composition is dispensed in a conventional aerosol can for shave gel compositions along with propellant.

30 35 The composition hereof can provide excellent shaving performance upon application to the face and shaving, along with

- 43 -

providing a long-lasting cooling sensation to the skin, with low skin sting.

Example 13

A skin lotion according to the present invention is exemplified below.

	<u>Ingredients (wt. %)</u>	<u>Example 13</u>
5	Carbomer 1	0.40
	Ethanol	10.00
	Glycerin	3.00
10	Perfume Component	0.80
	Sodium Hydroxide	0.15
	Coolant 1	0.18
	Coolant 2	0.07
	Water	q.s. to 100 %
15	1. Carbopol 954 and Pemulen TR 2 (5:3 weight ratio), both being homopolymers of acrylic acid crosslinked with an allyl ether of pentaerythritol or an allyl ether of sucrose, available from B. F. Goodrich Co. (Brecksville, OH, USA)	
20	The lotion is made by first mixing the water, carbomer, and glycerin. Next, a second mixture of the perfume component, ethanol, and coolants is prepared. Coolants 1 and 2 are as previously defined. Optionally, 0.25% of Coolant 3 as previously defined, can be added. The two mixtures are mixed together and neutralized with the NaOH. The resulting compositions can provide excellent skin moisturizing along with a long lasting cooling sensation with low skin sting.	
25		

Example 14

A skin lotion according to the present invention is exemplified below.

	<u>Ingredients (wt. %)</u>	<u>Example 14</u>
30	Carbomer 1	0.33
	Octyl Methoxycinnamate	7.50
	Triethanolamine	1.30
	Glycerin	1.00
35	Stearic Acid	1.00

- 44 -

Cetyl Alcohol	1.00
Cetyl Palmitate	0.50
DEA-Cetyl Phosphate ²	0.75
Perfume Component	0.25
5 Coolant 1	0.18
Coolant 2	0.07
Water	q.s. to 100%

1 Carbomer, as in Example 13.

2 Diethanolamide salt of cetyl phosphate

10 The sunscreen composition is prepared by mixing the water, carbomer, and glycerin and heating the mixture to about 80°C. The octyl methoxycinnamate, fatty alcohols, and stearic acid are then separately mixed together and heated to about 80°C. The two mixtures are mixed together and cooled to room temperature. The 15 coolants and perfume component are separately mixed and then added to the composition. Coolants 1 and 2 are as defined above. Optionally, 0.25% of Coolant 3, as defined above, can be included. Finally the triethanolamide is mixed into the composition.

20 The composition, upon application to the skin, can provide the skin with excellent protection from U.V. light while also providing a long-lasting cooling sensation without skin sting.

25

30

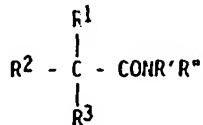
35

CA002491

- 45 -

WHAT IS CLAIMED IS:

1. A coolant composition for topical application to the skin characterized in that it comprises:
 (I) first coolant component which is an acyclic carboxamide coolant component of the formula:



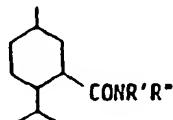
where:

(i) R' and R'' independently are hydrogen, C₁-C₅ alkyl, or C₁-C₈ hydroxyalkyl, R' and R'' provide a total of no more than 8 carbon atoms, and when R' is hydrogen, R'' may also be alkyl-carboxyalkyl of up to 6 carbon atoms

(ii) R' and R'' , when taken together, represent an alkylenes group of up to 6 carbon atoms, the opposite ends of which are attached to the amide nitrogen atom to form a nitrogen heterocycle, the carbon chain of which optionally being interrupted by oxygen;

(iii) R^1 is hydrogen or C₁-C₅ alkyl; R^2 and R^3 independently are C₁-C₅ alkyl; with the proviso that: (a) R^1 , R^2 , and R^3 together provide a total of at least 5 carbon atoms; and (b) when R^1 is hydrogen, R^2 is C₂-C₅ alkyl and R^3 is C₃-C₅ alkyl and at least one of R^2 and R^3 is branched;

- (II) a second coolant component which is a 3-substituted-p-methane of the formula:



where

R' is hydrogen or an aliphatic radical containing up to 25 carbon atoms;

R'' is hydroxy or an aliphatic radical containing up to 25 carbon atoms, with the proviso that when R' is hydrogen R'' may also be an aryl radical of up to 10 carbon atoms selected

- 46 -

from the group consisting of substituted phenyl, phenalkyl and substituted naphthyl, and pyridyl; and
R' and R", when taken together with the nitrogen atom to which they are attached, represent a cyclic or heterocyclic group of up to 25 carbon atoms;

(III) an aqueous vehicle; and

(IV) a cosmetically active ingredient or medicament, or a mixture thereof;

wherein said composition contains no more than 30% by weight of C₁-C₆ monohydric alcohol.

2. A coolant composition as in Claim 1 wherein said composition further comprises a perfume.

3. A coolant composition as in Claim 1, wherein the weight ratio of component (I) to component (II) is from 5:1 to 1:5, preferably from 3:1 to 1:3, more preferably from 3:1 to 1:1.

4. A coolant composition as in Claim 1, 2, or 3, wherein for said coolant component (I),

R' is hydrogen

R" is C₁-C₅ alkyl, C₁-C₈ hydroxyalkyl or alkylcarboxyalkyl of up to 6 carbon atoms;

R¹ is hydrogen or C₁-C₅ alkyl;

R² and R³ independently are C₁-C₅ alkyl;

R¹, R², and R³ together provide a total of at least 5 carbon atoms; and

when R¹ is hydrogen, R² is C₂-C₅ alkyl and R³ is C₃-C₅ alkyl branched at the carbon atom in the alpha or beta position.

5. A coolant composition as in Claim 4, wherein R¹ is methyl, ethyl or n-propyl, and at least one of R² and R³ has at least three carbon atoms and is branched at the alpha or beta carbon atom, and wherein R¹, R² and R³ preferably provide a total of from 5-10 carbon atoms.

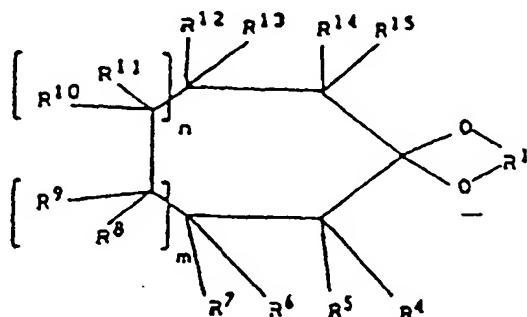
- 47 -

6. A coolant composition as in Claim 5, wherein component (I) is N,2,3-trimethyl-2-isopropyl butanamide.

7. A coolant composition as in Claim 1, 2, 3, 4, 5, or 6, wherein for said coolant component (II), R' and R" independently are hydrogen, C₁-C₆, straight or branched chain alkyl, C₁-C₆, straight or branched chain hydroxyalkyl or aminoalkyl or a C₁-C₆, acylated derivative thereof, or -C_nH_{2n}COR''' or -C_nH_{2n}COOR''' where -C_nH_{2n} is a straight or branched chain alkylene group in which n is an integer of from 1-6 and R''' is hydrogen or C₁-C₆ alkyl.

8. A coolant composition as in Claim 7, wherein said component (II) is N-ethyl-p-methane-3-carboxamide.

9. A coolant composition as in any of Claims 1-8, further comprising a coolant of the formula:



in which R⁴ to R¹⁵ independently of one another denote hydrogen or C₁-C₆ alkyl and m and n independently of one another denote zero or 1.

10. A coolant composition as in any of Claims 1-9 which contains no more than 25%, by weight, of C₁-C₆ monohydric alcohol, preferably no more than 10%, by weight, C₁-C₆ monohydric alcohol,

- 48 -

more preferably no more than 5% by weight, C₁-C₆ monohydric alcohol.

11. A coolant composition as in any of Claims 1-10, further comprising a cosmetically active ingredient selected from the group consisting of surfactants, conditioners, perfumes, and sunscreens, or a medicament selected from the group consisting of anti-acne and anti-inflammatory active ingredients, or a mixture thereof.

12. A method of providing a cooling sensation to the skin comprising applying an effective amount for providing a cooling sensation of the composition of any of Claims 1-11 to the skin.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 93/05455

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC Int.CI. 5 A61K7/00; A61K7/48; A61K7/15		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
Int.CI. 5	A61K	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
Y	US,A,4 136 163 (H. R. WATSON ET AL.) 23 January 1979 cited in the application see the whole document ---	1-8, 10-12
Y	US,A,4 230 688 (D.G. ROWSELL ET AL.) 28 October 1980 cited in the application see the whole document ---	1-8, 10-12
Y	LU,A,67 436 (WILKINSON SWORD LIMITED) 5 July 1973 cited in the application * whole document, especially examples VI,VII,X,XVII,XIX,XXV,XXVI,XXXVI * & US,A,4 230 688 ---	1-8, 10-12 -/-
<p>¹⁰ Special categories of cited documents : ¹¹ "A" document defining the general state of the art which is not considered to be of particular relevance ¹² "E" earlier document but published on or after the international filing date ¹³ "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) ¹⁴ "O" document referring to an oral disclosure, use, exhibition or other means ¹⁵ "P" document published prior to the international filing date but later than the priority date claimed ¹⁶ "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention ¹⁷ "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step ¹⁸ "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art ¹⁹ "Z" document member of the same patent family </p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search 07 OCTOBER 1993	Date of Mailing of this International Search Report 5. 10. 93	
International Searching Authority EUROPEAN PATENT OFFICE	Signature of Authorized Officer GAC G.	

Form PCT/ISA/250 (second sheet) (January 1985)

CA 002496

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
A	GB,A,2 233 873 (WARNER-LAMBERT COMPANY) 23 January 1991 see the whole document ---	1-8, 10-12
A	DE,A,2 413 639 (WILKINSON SWORD LTD) 17 October 1974 * whole document, especially examples 4,9,11,14,16 *	1,2,7,8, 10-12
A	JOURNAL OF THE SOCIETY OF COSMETIC CHEMISTS vol. 29, no. 4, April 1978, pages 185 - 200 WATSON ET AL. 'New compounds with the menthol cooling effect' see the whole document ---	1-8,12
A	FR,A,1 224 132 (OREC) 22 June 1960 see the whole document ---	9
P,A	EP,A,0 507 190 (HAARMANN & REIMER GMBH) 7 October 1992 see the whole document ----	9

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9305455
SA 75772

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 07/10/93

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
US-A-4136163	23-01-79	AT-B-	330361	25-06-76
		AU-B-	456393	29-11-74
		AU-A-	3860772	09-08-73
		BE-A-	778953	04-08-72
		CA-A-	969181	10-06-75
		CH-A-	556822	13-12-74
		CH-A-	548742	15-05-74
		DE-A,B,C	2205255	30-11-72
		FR-A,B	2127013	13-10-72
		GB-A-	1351761	01-05-74
		LU-A-	64729	04-07-72
		NL-A-	7201523	08-08-72
		SE-B-	397189	24-10-77
		US-A-	4193936	18-03-80
		US-A-	4178459	11-12-79
		US-A-	4226988	07-10-80
		US-A-	4150052	17-04-79
		US-A-	4190643	26-02-80
		AT-B-	335069	25-02-77
		AU-B-	457073	23-12-74
		AU-A-	3860572	09-08-73
		AU-B-	450078	07-06-74
		AU-A-	3860672	09-08-73
		BE-A-	778950	04-08-72
		BE-A-	778951	04-08-72
		CA-A-	968141	27-05-75
		CA-A-	1019551	25-10-77
		CH-A-	549952	14-06-74
		CH-A-	554648	15-10-74
		DE-A,B,C	2202535	17-08-72
		DE-A-	2203947	17-08-72
		FR-A,B	2127010	13-10-72
		FR-A,B	2127011	13-10-72
		GB-A-	1353381	15-05-74
		LU-A-	64713	30-06-72
		NL-A-	7201522	08-08-72
		NL-A-	7201524	08-08-72
		SE-B-	402404	03-07-78
		US-A-	4060091	29-11-77

EPO FORM P001

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

CA 002498

ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO. US 9305455
 SA 75772

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
 The members are as contained in the European Patent Office EDP file on
 The European Patent Office is in no way liable for those particulars which are merely given for the purpose of information. 07/10/93
 Page 2

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
US-A-4230688	28-10-80	GB-A-	1421743	21-01-76
		AT-B-	330963	26-07-76
		AU-B-	468442	15-01-76
		AU-A-	5442973	17-10-74
		BE-A-	798312	17-10-73
		CA-A-	984745	02-03-76
		CH-A-	564914	15-08-75
		DE-A,C	2317538	31-10-73
		DE-A,C	2366423	23-09-82
		FR-A,B	2180942	30-11-73
		JP-C-	1259130	12-04-85
		JP-A-	49019043	20-02-74
		JP-B-	59035940	31-08-84
		LU-A-	67436	05-07-73
		NL-A-	7305509	22-10-73
		SE-B-	418453	09-06-81
		US-A-	4296255	20-10-81
		US-A-	4153679	08-05-79
-----	-----	-----	-----	-----
LU-A-67436	05-07-73	GB-A-	1421743	21-01-76
		GB-A-	1411786	29-10-75
		AT-B-	330963	26-07-76
		AU-B-	468442	15-01-76
		AU-A-	5442973	17-10-74
		BE-A-	798312	17-10-73
		CA-A-	984745	02-03-76
		CH-A-	564914	15-08-75
		DE-A,C	2317538	31-10-73
		DE-A,C	2366423	23-09-82
		FR-A,B	2180942	30-11-73
		JP-C-	1259130	12-04-85
		JP-A-	49019043	20-02-74
		JP-B-	59035940	31-08-84
		NL-A-	7305509	22-10-73
		SE-B-	418453	09-06-81
		US-A-	4230688	28-10-80
		US-A-	4296255	20-10-81
		US-A-	4153679	08-05-79
		CA-A-	985168	09-03-76
		DE-A,C	2317539	31-10-73

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

EPO FORM P007

CA 002499

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9305455
SA 75772

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 07/10/93
Page 3

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
LU-A-67436		FR-A, B JP-C- JP-A- JP-B- NL-A-	2180943 1236697 49019044 59007744 7305508	30-11-73 31-10-84 20-02-74 20-02-84 22-10-73
GB-A-2233873	23-01-91	US-A- CA-A- JP-A-	5009893 2021222 3053849	23-04-91 18-01-91 07-03-91
DE-A-2413639	17-10-74	GB-A- CA-A- JP-A- NL-A-	1422998 1041506 50029753 7403797	28-01-76 31-10-78 25-03-75 24-09-74
FR-A-1224132		CH-A- DE-B- GB-A-	372134 1082370 850436	
EP-A-0507190	07-10-92	DE-A- AU-A- JP-A-	4110973 1390992 5125073	08-10-92 08-10-92 21-05-93

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82